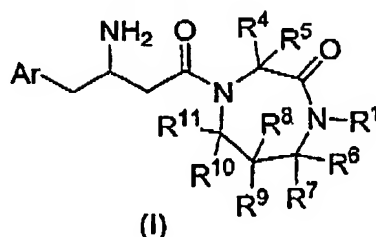


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**Listing of Claims:**

1. (original) A compound of the formula I:



or a pharmaceutically acceptable salt thereof; wherein  
 each n is independently 0, 1, or 2;

Ar is phenyl substituted with one to five  $R^3$  substituents;

$R^1$  is selected from the group consisting of  
 hydrogen,

$C_{1-10}$  alkyl, wherein alkyl is unsubstituted or substituted with one to five substituents  
 independently selected from halogen, hydroxy,  $C_{1-6}$  alkoxy, carboxy,  $C_{1-6}$   
 alkyloxycarbonyl, and phenyl- $C_{1-3}$  alkoxy, wherein alkoxy is unsubstituted or  
 substituted with one to five halogens,

$(CH_2)_n$ -aryl, wherein aryl is unsubstituted or substituted with one to five substituents  
 independently selected from halogen, CN, hydroxy,  $R^2$ ,  $OR^2$ ,  $NHSO_2R^2$ ,  
 $NR^2SO_2R^2$ ,  $SO_2R^2$ ,  $CO_2H$ , and  $C_{1-6}$  alkyloxycarbonyl,

$(CH_2)_n$ -heteroaryl, wherein heteroaryl is unsubstituted or substituted with one to three  
 substituents independently selected from hydroxy, halogen,  $C_{1-6}$  alkyl, and  $C_{1-6}$   
 alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five  
 halogens,

$(CH_2)_n$ -heterocyclyl, wherein heterocyclyl is unsubstituted or substituted with one to  
 three substituents independently selected from oxo, hydroxy, halogen,  $C_{1-6}$  alkyl,  
 and  $C_{1-6}$  alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with  
 one to five halogens,

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(CH<sub>2</sub>)<sub>n</sub>-C<sub>3-6</sub> cycloalkyl, wherein cycloalkyl is unsubstituted or substituted with one to three substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; and  
wherein any methylene (CH<sub>2</sub>) carbon atom in (CH<sub>2</sub>)<sub>n</sub> is unsubstituted or substituted with one to two groups independently selected from halogen, hydroxy, and C<sub>1-4</sub> alkyl unsubstituted or substituted with one to five halogens;

each R<sup>3</sup> is independently selected from the group consisting of  
hydrogen,  
halogen,  
cyano,  
hydroxy,  
C<sub>1-6</sub> alkyl, unsubstituted or substituted with one to five halogens,  
C<sub>1-6</sub> alkoxy, unsubstituted or substituted with one to five halogens,  
carboxy,  
alkoxycarbonyl,  
amino,  
NHR<sup>2</sup>,  
NR<sup>2</sup>R<sup>2</sup>,  
NHSO<sub>2</sub>R<sup>2</sup>,  
NR<sup>2</sup>SO<sub>2</sub>R<sup>2</sup>,  
NHCOR<sup>2</sup>,  
NR<sup>2</sup>COR<sup>2</sup>,  
NHCO<sub>2</sub>R<sup>2</sup>,  
NR<sup>2</sup>CO<sub>2</sub>R<sup>2</sup>,  
SO<sub>2</sub>R<sup>2</sup>,  
SO<sub>2</sub>NH<sub>2</sub>,  
SO<sub>2</sub>NHR<sup>2</sup>, and  
SO<sub>2</sub>NR<sup>2</sup>R<sup>2</sup>;

each R<sup>2</sup> is independently C<sub>1-6</sub> alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, CO<sub>2</sub>H, and C<sub>1-6</sub> alkyloxycarbonyl;

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R<sup>4</sup>, R<sup>6</sup>, and R<sup>10</sup> are each independently selected from the group consisting of:

hydrogen,

cyano,

carboxy,

C<sub>1-6</sub> alkyloxycarbonyl,

C<sub>1-10</sub> alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkoxy, carboxy,

C<sub>1-6</sub> alkyloxycarbonyl, and phenyl-C<sub>1-3</sub> alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens,

(CH<sub>2</sub>)<sub>n</sub>-aryl, wherein aryl is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,

(CH<sub>2</sub>)<sub>n</sub>-heteroaryl, wherein heteroaryl is unsubstituted or substituted with one to three substituents independently selected from hydroxy, halogen, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,

(CH<sub>2</sub>)<sub>n</sub>-heterocyclyl, wherein heterocyclyl is unsubstituted or substituted with one to three substituents independently selected from oxo, hydroxy, halogen, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,

(CH<sub>2</sub>)<sub>n</sub>-C<sub>3-6</sub> cycloalkyl, wherein cycloalkyl is unsubstituted or substituted with one to three substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,

(CH<sub>2</sub>)<sub>n</sub>CONR<sup>12</sup>R<sup>13</sup>, wherein R<sup>12</sup> and R<sup>13</sup> are independently selected from the group consisting of hydrogen, tetrazolyl, thiazolyl, (CH<sub>2</sub>)<sub>n</sub>-phenyl, (CH<sub>2</sub>)<sub>n</sub>-C<sub>3-6</sub> cycloalkyl, and C<sub>1-6</sub> alkyl, wherein alkyl is unsubstituted or substituted with one to five halogens and wherein phenyl and cycloalkyl are unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; or wherein R<sup>12</sup> and R<sup>13</sup> together with the nitrogen atom to which they are attached form a heterocyclic ring selected from azetidine, pyrrolidine, piperidine, piperazine, and morpholine wherein said

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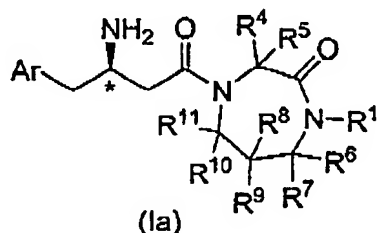
heterocyclic ring is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens;  
 and wherein any methylene (CH<sub>2</sub>) carbon atom in (CH<sub>2</sub>)<sub>n</sub> is unsubstituted or substituted with one to two groups independently selected from halogen, hydroxy, and C<sub>1-4</sub> alkyl unsubstituted or substituted with one to five halogens;

R<sup>8</sup> is selected from the group consisting of halogen, hydroxy, and R<sup>4</sup>;

R<sup>5</sup>, R<sup>7</sup> and R<sup>11</sup> are each independently hydrogen or C<sub>1-6</sub> alkyl; or wherein R<sup>7</sup> and R<sup>1</sup> together with the nitrogen atom to which R<sup>1</sup> is attached form a heterocyclic ring selected from azetidine, pyrrolidine and piperidine wherein said heterocyclic ring is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; and

R<sup>9</sup> is selected from the group consisting of hydrogen, hydroxy, halogen, or C<sub>1-6</sub> alkyl; with the proviso that at least one of R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> is not hydrogen.

2. (original) The compound of Claim 1 of the formula Ia:



wherein the carbon atom marked with an \* has the *R* configuration.

3. (original) The compound of Claim 1 wherein R<sup>3</sup> is selected from the group consisting of hydrogen, fluoro, chloro, bromo, trifluoromethyl, and methyl.

4. (original) The compound of Claim 3 wherein R<sup>3</sup> is hydrogen, chloro, or fluoro.

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5. (original) The compound of Claim 1 wherein  $R^1$  is selected from the group consisting of  
hydrogen,  
 $C_{1-6}$  alkyl, wherein alkyl is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy,  $C_{1-6}$  alkoxy, carboxy,  $C_{1-6}$  alkyloxycarbonyl, and phenyl- $C_{1-3}$  alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens, and  
 $(CH_2)_n$ - $C_{3-6}$  cycloalkyl, wherein cycloalkyl is unsubstituted or substituted with one to three substituents independently selected from halogen, hydroxy,  $C_{1-6}$  alkyl, and  $C_{1-6}$  alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; and  
wherein any methylene  $(CH_2)$  carbon atom in  $(CH_2)_n$  is unsubstituted or substituted with one to two groups independently selected from halogen, hydroxy, and  $C_{1-4}$  alkyl unsubstituted or substituted with one to five halogens.

6. (original) The compound of Claim 5 wherein  $R^1$  is selected from the group consisting of hydrogen, methyl, and cyclopropyl.

7. (original) The compound of Claim 6 wherein  $R^1$  is hydrogen.

8. (original) The compound of Claim 1 wherein  $R^4$  is selected from the group consisting of:

hydrogen,  
 $C_{1-6}$  alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy,  $C_{1-6}$  alkoxy, carboxy,  $C_{1-6}$  alkyloxycarbonyl, and phenyl- $C_{1-3}$  alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens,  
 $(CH_2)_n$ -aryl, wherein aryl is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy,  $C_{1-6}$  alkyl, and  $C_{1-6}$  alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,  
 $(CH_2)_n$ -heteroaryl, wherein heteroaryl is unsubstituted or substituted with one to three substituents independently selected from hydroxy, halogen,  $C_{1-6}$  alkyl, and  $C_{1-6}$

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alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,  
(CH<sub>2</sub>)<sub>n</sub>-C<sub>3-6</sub> cycloalkyl, wherein cycloalkyl is unsubstituted or substituted with one to three substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; and  
wherein any methylene (CH<sub>2</sub>) carbon atom in (CH<sub>2</sub>)<sub>n</sub> is unsubstituted or substituted with one to two groups independently selected from halogen, hydroxy, and C<sub>1-4</sub> alkyl unsubstituted or substituted with one to five halogens.

9. (original) The compound of Claim 8 wherein R<sup>4</sup> is selected from the group consisting of:

hydrogen,  
CH<sub>3</sub>,  
CH<sub>2</sub>CH<sub>3</sub>,  
CH<sub>2</sub>CF<sub>3</sub>,  
CH<sub>2</sub>(2-pyridyl),  
CH<sub>2</sub>Ph,  
CH<sub>2</sub>(2-F-Ph),  
CH<sub>2</sub>(2-Me-Ph), and  
CH<sub>2</sub>(2-CF<sub>3</sub>-Ph).

10. (original) The compound of Claim 1 wherein R<sup>6</sup> is selected from the group consisting of:

hydrogen,  
C<sub>1-6</sub> alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkoxy, carboxy, C<sub>1-6</sub> alkyloxycarbonyl, and phenyl-C<sub>1-3</sub> alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens,  
(CH<sub>2</sub>)<sub>n</sub>-aryl, wherein aryl is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,  
(CH<sub>2</sub>)<sub>n</sub>-heteroaryl, wherein heteroaryl is unsubstituted or substituted with one to three substituents independently selected from hydroxy, halogen, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub>

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alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens,  
(CH<sub>2</sub>)<sub>n</sub>-C<sub>3-6</sub> cycloalkyl, wherein cycloalkyl is unsubstituted or substituted with one to three substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; and  
wherein any methylene (CH<sub>2</sub>) carbon atom in (CH<sub>2</sub>)<sub>n</sub> is unsubstituted or substituted with one to two groups independently selected from halogen, hydroxy, and C<sub>1-4</sub> alkyl unsubstituted or substituted with one to five halogens.

11. (original) The compound of Claim 10 wherein R<sup>6</sup> is selected from the group consisting of:

hydrogen,  
C<sub>1-6</sub> alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkoxy, carboxy, C<sub>1-6</sub> alkyloxycarbonyl, and phenyl-C<sub>1-3</sub> alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens, and  
(CH<sub>2</sub>)<sub>n</sub>-aryl, wherein aryl is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens; and  
wherein methylene (CH<sub>2</sub>) carbon atom in (CH<sub>2</sub>)<sub>n</sub> is unsubstituted or substituted with one to two groups independently selected from halogen, hydroxy, and C<sub>1-4</sub> alkyl unsubstituted or substituted with one to five halogens.

12. (original) The compound of Claim 11 wherein R<sup>6</sup> is selected from the group consisting of:

hydrogen,  
CH<sub>3</sub>,  
CH<sub>2</sub>CH<sub>3</sub>,  
CF<sub>3</sub>,  
CH<sub>2</sub>Ph, and  
CH<sub>2</sub>(2-F-Ph).

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13. (original) The compound of Claim 1 wherein R<sup>8</sup> is selected from the group consisting of:

hydrogen,

hydroxy,

halogen, and

C<sub>1-6</sub> alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkoxy, carboxy, C<sub>1-6</sub> alkyloxycarbonyl, and phenyl-C<sub>1-3</sub> alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens.

14. (original) The compound of Claim 13 wherein R<sup>8</sup> is hydrogen.

15. (original) The compound of Claim 1 wherein R<sup>10</sup> is selected from the group consisting of:

hydrogen, and

C<sub>1-6</sub> alkyl, unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkoxy, carboxy, C<sub>1-6</sub> alkyloxycarbonyl, and phenyl-C<sub>1-3</sub> alkoxy, wherein alkoxy is unsubstituted or substituted with one to five halogens.

16. (original) The compound of Claim 15 wherein R<sup>10</sup> is hydrogen.

17. (original) The compound of Claim 1 wherein R<sup>5</sup>, R<sup>7</sup> and R<sup>11</sup> are each independently selected from hydrogen and methyl.

18. (original) The compound of Claim 17 wherein R<sup>5</sup>, R<sup>7</sup> and R<sup>11</sup> are hydrogen.

19. (original) The compound of Claim 1 wherein R<sup>9</sup> is selected from hydrogen, halogen and methyl.

20. (original) The compound of Claim 19 wherein R<sup>9</sup> is hydrogen.

21. (original) The compound of Claim 19 wherein R<sup>9</sup> is methyl and R<sup>5</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>10</sup>, and R<sup>11</sup> are hydrogen.



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22. (original) The compound of Claim 21 wherein R<sup>4</sup> is selected from the group consisting of:

hydrogen,  
CH<sub>3</sub>,  
CH<sub>2</sub>CH<sub>3</sub>,  
CH<sub>2</sub>CF<sub>3</sub>,  
CH<sub>2</sub>(2-pyridyl),  
CH<sub>2</sub>Ph,  
CH<sub>2</sub>(2-F-Ph),  
CH<sub>2</sub>(2-Me-Ph), and  
CH<sub>2</sub>(2-CF<sub>3</sub>-Ph).

23. (original) The compound of Claim 1 wherein R<sup>5</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, and R<sup>11</sup> are hydrogen, with the proviso that R<sup>6</sup> is not hydrogen.

24. (original) The compound of Claim 23 wherein R<sup>4</sup> is selected from the group consisting of:

hydrogen,  
CH<sub>3</sub>,  
CH<sub>2</sub>CH<sub>3</sub>,  
CH<sub>2</sub>CF<sub>3</sub>,  
CH<sub>2</sub>(2-pyridyl),  
CH<sub>2</sub>Ph,  
CH<sub>2</sub>(2-F-Ph),  
CH<sub>2</sub>(2-Me-Ph), and  
CH<sub>2</sub>(2-CF<sub>3</sub>-Ph); and

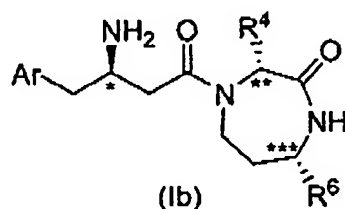
R<sup>6</sup> is selected from the group consisting of:

CH<sub>3</sub>,  
CH<sub>2</sub>CH<sub>3</sub>,  
CF<sub>3</sub>,  
CH<sub>2</sub>Ph, and  
CH<sub>2</sub>(2-F-Ph).

25. (original) The compound of Claim 24 wherein R<sup>1</sup> is hydrogen.

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26. (original) The compound of Claim 25 wherein the stereogenic carbon atoms marked with an \*\* and an \*\*\* have the stereochemistry as depicted in formula Ib:



27. (original) The compound of Claim 1 wherein R<sup>7</sup> and R<sup>1</sup> together with the nitrogen atom to which R<sup>1</sup> is attached form a heterocyclic ring selected from azetidine, pyrrolidine and piperidine wherein said heterocyclic ring is unsubstituted or substituted with one to five substituents independently selected from halogen, hydroxy, C<sub>1-6</sub> alkyl, and C<sub>1-6</sub> alkoxy, wherein alkyl and alkoxy are unsubstituted or substituted with one to five halogens.

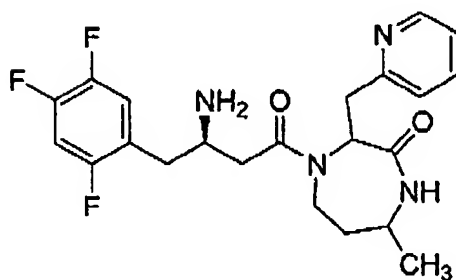
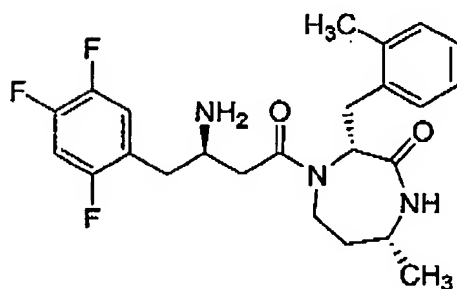
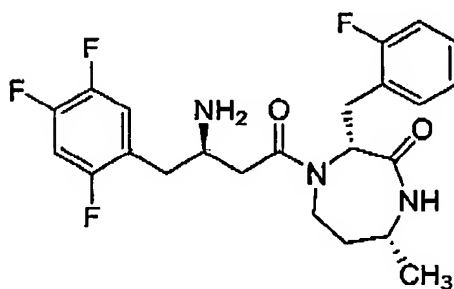
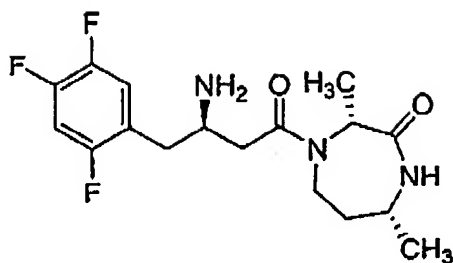
28. (original) The compound of Claim 27 wherein R<sup>7</sup> and R<sup>1</sup> together with the nitrogen atom to which R<sup>1</sup> is attached form a pyrrolidine ring.

29. (original) The compound of Claim 28 wherein R<sup>4</sup> is selected from the group consisting of:

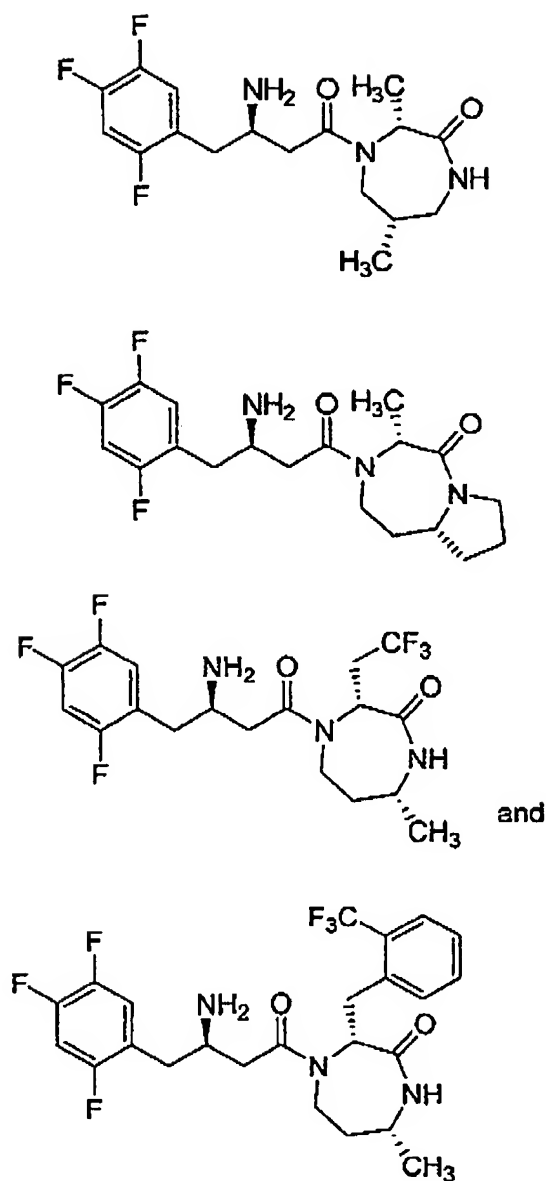
hydrogen,  
 CH<sub>3</sub>,  
 CH<sub>2</sub>CH<sub>3</sub>,  
 CH<sub>2</sub>CF<sub>3</sub>,  
 CH<sub>2</sub>(2-pyridyl),  
 CH<sub>2</sub>Ph,  
 CH<sub>2</sub>(2-F-Ph),  
 CH<sub>2</sub>(2-Me-Ph), and  
 CH<sub>2</sub>(2-CF<sub>3</sub>-Ph).

30. (original) A compound selected from the group consisting of:

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or a pharmaceutically acceptable salt thereof.

31. (original) A pharmaceutical composition which comprises a compound of Claim 1 and a pharmaceutically acceptable carrier.

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32. (previously amended) A method of treating Type 2 diabetes in a mammal in need thereof which comprises the administration to the mammal of a therapeutically effective amount of a compound of Claim 1.

33-34. (previously cancelled)